

AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS:

1.-12. (canceled)

13. (withdrawn-currently amended): A method of preparing thea *Mycobacterium* promoter of claim 23 expression system for high throughput screening and developing inhibitors against mycobacteria under low carbon source, said process comprising the steps of:

(a) isolating said promoter from *Mycobacterium* DNA and characterizing a 200 bp promoter sequence having SEQ ID NO. 2 from nucleotide sequence of 1.5 kb DNA fragment upstream of *M. tuberculosis* gene *relA/spoT*,

(b) ligating the isolated promoter sequence of step (a) into a plasmid vector vector pSAK12, and

(c) studying the expression of the promoter sequence under low carbon source or carbon starved conditions.

14. (withdrawn-currently amended): The A process of as claimed in claim 13, wherein the *Mycobacterium* promoter is 2.5 foldmore active in *M. Smegmatis* than the conventional heat shock protein promoter (P_{hsp60})(heat shock protein promoter) promoter.

15. (withdrawn-currently amended): A process of expressing a reporter gene in *M. smegmatis* under carbon starved conditions, the process comprising the step of growing *M. smegmatis* containing the promoter of claim 28 as claimed in claim 13, wherein the carbon source is about 2.5 to 0.001% glucose is in the range of about 2.5 to 0.001%.

16. (withdrawn-currently amended): TheA process of as claimed in claim 1514, wherein the carbon source is about 2 to 0.02%, glucose is in the range of about 2 to 0.02%.

17. (withdrawn-currently amended): TheA process of as claimed in claim 1513, wherein the percentage inhibition growth of the *M. smegmatis* mycobacteria in presence of the promoter and in presence of inhibitor ethambutol is reduced by in the range of about 6 to 25% by the presence of ethambutol in presence of 0.02 % glucose i.e under starved conditions.

18. (withdrawn-currently amended): TheA process of as claimed in claim 17, wherein the percentage inhibition growth of the *M. smegmatis* mycobacteria in the presence of the promoter and in presence of inhibitor ethambutol is reduced by in the range of about 7 to 21% by the presence of ethambutol in presence of 0.02 % glucose i.e under starved conditions.

19. (withdrawn-currently amended): TheA process of as claimed in claim 1513, wherein the percentage inhibition growth of the *M. smegmatis* mycobacteria in presence of the promoter

and in presence of inhibitor isoniazide is reduced by in the range of about 15 to 45% by the presence of isoniazid in presence of 0.02 % glucose i.e under starved conditions.

20. (withdrawn-currently amended): TheA process ofas claimed in claim 19, wherein thepercentage inhibition growth of the *M. smegmatis* mycobacteria in presence of the promoter and in presence of inhibitor isoniazide is reduced by in the range of about 18 to 40 % in the presence of isoniazid in presence of 0.02 % glucose i.e under starved conditions.

21. (withdrawn-currently amended): TheA process ofas claimed in claim 1513, wherein thepercentage inhibition growth of the *M. smegmatis* mycobacteria in presence of the promoter and in presence of inhibitor rifampicin is reduced by in the range of about 20 to 45% by the presence of rifampicin in presence of 0.02 % glucose i.e under starved conditions.

22. (withdrawn-currently amended): TheA process ofas claimed in claim 21, wherein thepercentage inhibition growth of the *M. smegmatis* mycobacteria in presence of the promoter and in presence of inhibitor rifampicin is reduced by in the range of about 21 to 41% by the presence of rifampicin in presence of 0.02 % glucose i.e under starved conditions.

23. (new) A *Mycobacterium* promoter, wherein the promoter is stable in *M. smegmatis* and *E. coli*, and consists essentially of the 200 base pair fragment upstream and adjacent to the *Mycobacterium tuberculosis* *relA/SpoT* gene.

24. (new) The *Mycobacterium* promoter of claim 23, wherein the promoter is operatively linked to a reporter gene.

25. (new) The *Mycobacterium* promoter of claim 24, wherein said reporter gene is LacZ.

26. (new) The *Mycobacterium* promoter of claim 24, wherein said reporter gene is xyle.

27. (new) The *Mycobacterium* promoter of claim 24, wherein the promoter is 2.5 fold more active in *M. smegmatis* than the heat shock protein promoter (P_{hsp60}).

28 (new) The *Mycobacterium* promoter of claim 24, wherein the promoter is further contained in a plasmid with an Ampicillin or Kanamycin resistance marker.

29. (new) The *Mycobacterium* promoter of claim 23, wherein the promoter consists of SEQ ID NO:2.